Amendments to the Claims

1. (previously presented) A compound represented by formula (I);

wherein:

R and R' are independently C₁-C₅ alkyl, or together R and R' form a saturated or unsaturated carbocyclic ring having from 3 to 8 carbon atoms:

R_{PH} is hydrogen or methyl;

R1 and R2 are independently hydrogen, halo, or C1-C5 alkyl;

L₁ is -(CH₂)_m-O-;

 L_2 is $-(CH_2)_mCH(OH)$ - or $-(CH_2)_mC(O)$ -;

where m is 0, 1 or 2,

RB is a branched C3-C5 alkyl,

RC is:

-O-SO₂-(R50) where R50 is -C₁₋₃alkyl, or -(CH₂)₁₋₂CF₃.

-NH-SO₂-(R50), where R50 is -C₁₋₃alkyl, -CF₃, or -(CH₂)₁₋₂CF₃;

-N(CH $_3$)-SO $_2$ -C $_{1-2}$ alkyl; or

-N(SO₂R51)₂ where each R51 is independently, -C₁₋₃alkyl, -CF₃, or -(CH₂)₁₋₂CF₃.

2. (previously presented) A compound according to Claim 1 wherein R_{PH} is hydrogen.

3-4. (canceled)

5. (previously presented) A compound according to Claim 1 represented by the structural formulae below as follows:

M-1)

M-2)

M-3)

M-6)

M-7)

M-8)

M-9)

M-11)

M-13)

M-14)

M-15)

M-16)

M-17)

M-18)

M-19)

M-20)

6. (previously presented) A compound represented by the structural formulae M-32 to M-50 as follows:

M-32)

M-34)

M-35)

M-36)

M-37)

M-38)

M-39)

M-40)

M-41)

M-42)

M-43)

M-44)

M-45)

M-46)

M-47)

M-48)

M-49)

M-50)

7. (canceled)

8. (original) A compound represented by a formula below:

9-15. (canceled)

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16. (previously presented) A pharmaceutical formulation comprising the compound according to Claim 1 together with a pharmaceutically acceptable carrier or diluent.

17-18. (canceled)

19. (original) A formulation for treating psoriasis comprising:

Ingredient (A2): the vitamin D receptor modulator of claim 1:

Ingredient (B2):

one or more co-agents that are conventional for treatment psoriasis selected from the group consisting of:

a. topical glucocorticoids.

salicylic acid.

crude coal tar; and

Ingredient (C2): optionally, a carrier or diluent.

20. (canceled)

- 21. (previously presented) A method of treating a mammal for Osteoporosis, Psoriasis, Scleroderma, or seborrheic dermatitis wherein the method comprises administering a pharmaceutically effective amount of at least one compound of claim 1.
 - 22. (original) The method of claim 21 for the treatment of psoriasis.
 - 23. (original) The method of claim 21 for the treatment of osteoporosis.

24-32. (canceled)

33. (previously presented) A compound represented by formula (II) or a pharmaceutically acceptable salt thereof:

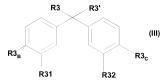
wherein:

R2 and R2' are independently methyl or ethyl;

R21 and R22 are independently selected from: hydrogen, methyl, ethyl, or -Cl, R2B is 3,3-dimethyl-2-hydroxybutoxy or 3,3-dimethyl-2-oxobutoxy; and R2C is

where Q is -O- or -NH-.

34. (previously presented) A compound represented by formula (III) or a pharmaceutically acceptable salt thereof:



wherein:

R3 and R3' are independently methyl or ethyl;

R31 and R32 are independently selected from: hydrogen, methyl, ethyl, or -Cl, R3_B is 3,3-dimethyl-2-hydroxybutoxy or 3,3-dimethyl-2-oxobutoxy; and

R_{3c} is

35. (previously presented) A compound represented by a formula below:

36. (previously presented) A compound represented by a formula below:

37. (previously presented) A compound represented by a formula below:

- 38. (previously presented) A pharmaceutical formulation comprising the compound according to one of claims 35, 36, or 37 together with a pharmaceutically acceptable carrier or diluent.
- 39. (previously presented) A method of treating a mammal for Osteoporosis, Psoriasis, Scleroderma, or seborrheic dermatitis wherein the method comprises administering a pharmaceutically effective amount of at least the compound of according to one of claims 35, 36, or 37.
 - 40. (previously presented) The method of claim 39 for the treatment of psoriasis.
 - 41. (previously presented) The method of claim 39 for the treatment of osteoporosis.